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Examiner Initials	Document Description
DMY MAY 15 2002	Kimberling et al., "Gene mapping of Usher syndrome type IIa: localization of the gene to a 2.1-cM segment on chromosome 1q41," <i>Am J Hum Genet.</i> 1995 Jan; 56(1):216-23.
AM	Leonardo et al., "Guidance of developing axons by netrin-1 and its receptors," <i>Cold Spring Harb Symp Quant Biol.</i> 1997;62:467-78.
	Lin, "Immunogold localization of extracellular matrix molecules in Bruch's membrane of the rat," <i>Curr Eye Res.</i> 1989 Nov;8(11):1171-8.
	Lindenov, <i>The Etiology of Deaf-mutism with Special Reference to Heredity</i> , Vol. 8 in series <i>Opera ex Domo Biologiae Hereditariae Humanae Universitatis Havnensis</i> ; Einar Munksgaard, Copenhagen, Denmark (1945) 6 pgs.
	Liu et al., "A mutation (2314delG) in the Usher syndrome type IIa gene: high prevalence and phenotypic variation," <i>Am J Hum Genet.</i> 1999 Apr;64(4):1221-5.
	Mattson et al., "A practical approach to crosslinking," <i>Mol Biol Rep.</i> 1993 Apr; 17(3):167-83.
	Mayer et al., "Low nitrogen affinity of laminin-5 can be attributed to two serine residues in EGF-like motif $\gamma 2$ III4," <i>FEBS Lett.</i> 1995 May 29;365(2-3): 129-32.
	Métin et al., "A role for netrin-1 in the guidance of cortical efferents," <i>Development.</i> 1997 Dec;124(24):5063-74.
M	National Center for Biotechnology Information, National Library of Medicine, National Institutes of Health, GenBank Locus AF055580, Accession No. AF055580, "Homo sapiens Usher syndrome type IIa protein (USH2A) mRNA, complete cds.," [online]. Bethesda, MD [retrieved on 2002-04-05]. Retrieved from the Internet: <URL: <a href="http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=2&amp;db=nucleotide&amp;page=0&amp;dispmax=20&amp;WebEnv=IEQSF%3EFJ%3FNIBNHg%3CB%3CCEcKAI%3E%3E%3F%3EfoICA%3CB%5D%3E%40%5CHJAkWcC%60I%3Ed%3C&amp;WebEnvRq=1">http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=2&amp;db=nucleotide&amp;page=0&amp;dispmax=20&amp;WebEnv=IEQSF%3EFJ%3FNIBNHg%3CB%3CCEcKAI%3E%3E%3F%3EfoICA%3CB%5D%3E%40%5CHJAkWcC%60I%3Ed%3C&amp;WebEnvRq=1</a> >; 4 pgs.
M	National Center for Biotechnology Information, National Library of Medicine, National Institutes of Health, GenBank Locus HSUSH2A01, Accession No. AF091873, "Homo sapiens Usher syndrome type IIa protein gene, exons 1 and 2," [online]. Bethesda, MD [retrieved on 2002-04-05]. Retrieved from the Internet: <URL: <a href="http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=4&amp;db=nucleotide&amp;page=0&amp;dispmax=20&amp;WebEnv=Wg%3Cn_FDG%5DE%60%3E%3D%3Cc%5DPGDJc_gTB%5EjbFkl%3C_JEH%3Dzcc%3EF%5EeffJdTI%3D%3F%3C&amp;WebEnvRq=1">http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=4&amp;db=nucleotide&amp;page=0&amp;dispmax=20&amp;WebEnv=Wg%3Cn_FDG%5DE%60%3E%3D%3Cc%5DPGDJc_gTB%5EjbFkl%3C_JEH%3Dzcc%3EF%5EeffJdTI%3D%3F%3C&amp;WebEnvRq=1</a> >; 3 pgs.

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